ALLERGY RELIEF NON DROWSY- loratadine tablet Bryant Ranch Prepack

Drug Facts

Active ingredient (in each tablet)

Loratadine, USP 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

Warnings

Do not use

if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product,

do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if

an allergic reaction to this product occurs. Seek medical help right away

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

adults and children 6 years and over 1 tablet daily; not more than 1 tablet in 24 hours children under 6 years of age ask a doctor consumers with liver or kidney disease ask a doctor

Other information

- store at 20°-25°C (68°-77°F) (see UPS Controlled Room Temperature)
- protect from light

Inactive ingredients

lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or comments?

Call 1-888-588-1418 Monday-Friday 9AM-5PM EST

HOW SUPPLIED

NDC: 71335-1651-1: 20 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-2: 30 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-3: 60 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-4: 14 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-5: 10 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-6: 90 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-6: 28 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-8: 15 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-8: 15 Tablets in a BOTTLE, PLASTIC NDC: 71335-1651-9: 100 Tablets in a BOTTLE, PLASTIC

Loratadine 10mg Tablet

Each tablet contains: Loratadine, USP 10 mg.

Keep this and all drugs out of the reach of children.

Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature].

Protect from light.

NDC 71335-1651-1

Loratadine Tablets, USP

10 mg

BRP

20 Tablets

Repackaged by: Bryant Ranch Prepack, Inc. Burbank, CA 91504 USA Manufactured by: Camber Consumer Care



ALLERGY RELIEF NON DROWSY

loratadine tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71335-1651(NDC:69230-317)

Route of Administration ORAL

Active Ingredient/Active Moiety

SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)

Ingredient Name
Basis of Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)
LORATADINE
10 mg

Inactive Ingredients Ingredient Name Strength LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X) MAGNESIUM STEARATE (UNII: 70097M6I30) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)

Product Characteristics

Color white Score no score

Shape ROUND Size 6mm

Flavor Imprint Code 439

Contains

| Packaging | | | | | | | |
|-----------|----------------------|---|-------------------------|-----------------------|--|--|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | | | |
| 1 | NDC:71335- 1651-1 | 20 in 1 BOTTLE; Type 0: Not a Combination Product | 07/27/2020 | | | | |
| ٦ | NDC:71335- | 30 in 1 BOTTLE; Type 0: Not a Combination | 07/16/2020 | | | | |

| | 1651-2 | Product | 0//10/2020 | |
|---|----------------------|--|------------|--|
| 3 | NDC:71335- 1651-3 | 60 in 1 BOTTLE; Type 0: Not a Combination Product | 10/16/2020 | |
| 4 | NDC:71335- 1651-4 | 14 in 1 BOTTLE; Type 0: Not a Combination Product | 05/25/2022 | |
| 5 | NDC:71335- 1651-5 | 10 in 1 BOTTLE; Type 0: Not a Combination Product | 01/12/2021 | |
| 6 | NDC:71335- 1651-6 | 90 in 1 BOTTLE; Type 0: Not a Combination Product | 07/01/2020 | |
| 7 | NDC:71335- 1651-7 | 28 in 1 BOTTLE; Type 0: Not a Combination Product | 04/04/2024 | |
| 8 | NDC:71335- 1651-8 | 15 in 1 BOTTLE; Type 0: Not a Combination Product | 04/04/2024 | |
| 9 | NDC:71335- 1651-9 | 100 in 1 BOTTLE; Type 0: Not a Combination Product | 01/14/2021 | |

| Marketing Information | | | | | | |
|-----------------------|---|-------------------------|-----------------------|--|--|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | | | |
| ANDA | ANDA075209 | 12/27/2019 | | | | |
| | | | | | | |

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

| Establishment | | | | | | |
|----------------------|---------|-----------|---|--|--|--|
| Name | Address | ID/FEI | Business Operations | | | |
| Bryant Ranch Prepack | | 171714327 | REPACK(71335-1651), RELABEL(71335-1651) | | | |

Revised: 4/2024 Bryant Ranch Prepack